

FORM PTO-1390 (Modified)  
(REV 11-2000)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

TRANSMITTAL LETTER TO THE UNITED STATES  
DESIGNATED/ELECTED OFFICE (DO/EO/US)  
CONCERNING A FILING UNDER 35 U.S.C. 371

A-7419

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 1.5)

09/831355

INTERNATIONAL APPLICATION NO.  
PCT/AU99/00795INTERNATIONAL FILING DATE  
20 September 1999 (20.09.99)PRIORITY DATE CLAIMED  
21 September 1998 (21.09.98)

TITLE OF INVENTION

METHOD FOR TREATING PRECIOUS METAL BEARING MINERALS

APPLICANT(S) FOR DO/EO/US

Michael HOURN; Rodrigo VENTURA; John WILLIS and David WINBORNE

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (24) indicated below.
4. ☐ The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371 (c) (2))
  - a. ☒ is attached hereto (required only if not communicated by the International Bureau).
  - b. ☐ has been communicated by the International Bureau.
  - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).
  - a. ☐ is attached hereto.
  - b. ☐ has been previously submitted under 35 U.S.C. 154(d)(4).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3))
  - a. ☐ are attached hereto (required only if not communicated by the International Bureau).
  - b. ☐ have been communicated by the International Bureau.
  - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired
  - d. ☒ have not been made and will not be made.
8. ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).
10. ☐ An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)).
11. ☒ A copy of the International Preliminary Examination Report (PCT/IPEA/409).
12. ☒ A copy of the International Search Report (PCT/ISA/210).

## Items 13 to 20 below concern document(s) or information included:

13. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
14. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
15. ☒ A **FIRST** preliminary amendment.
16. ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
17. ☐ A substitute specification.
18. ☐ A change of power of attorney and/or address letter.
19. ☐ A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
20. ☐ A second copy of the published international application under 35 U.S.C. 154(d)(4).
21. ☐ A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
22. ☐ Certificate of Mailing by Express Mail
23. ☐ Other items or information:

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 1.5) <b>09/831355</b>	INTERNATIONAL APPLICATION NO. <b>PCT/AU99/00795</b>	ATTORNEY'S DOCKET NUMBER <b>A-7419</b>
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24. The following fees are submitted:

**BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :**

- ☒ Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO . . . . . **\$1000.00**
- ☐ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO . . . . . **\$860.00**
- ☐ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO . . . . . **\$710.00**
- ☐ International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) . . . . . **\$690.00**
- ☐ International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) . . . . . **\$100.00**

**ENTER APPROPRIATE BASIC FEE AMOUNT =****CALCULATIONS PTO USE ONLY****\$1,000.00**

Surcharge of **\$130.00** for furnishing the oath or declaration later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).

**\$0.00**

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE
Total claims	24 - 20 =	4	x \$18.00
Independent claims	3 - 3 =	0	x \$80.00

**\$72.00****\$0.00**Multiple Dependent Claims (check if applicable). ☐**\$0.00****TOTAL OF ABOVE CALCULATIONS =****\$1,072.00**

Applicant claims small entity status. (See 37 CFR 1.27). The fees indicated above are reduced by 1/2.

**\$0.00****SUBTOTAL =****\$1,072.00**

Processing fee of **\$130.00** for furnishing the English translation later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).

**\$0.00****TOTAL NATIONAL FEE =****\$1,072.00**

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable). ☐

**\$0.00****TOTAL FEES ENCLOSED =****\$1,072.00**

Amount to be refunded	\$
charged	\$

- a. ☒ A check in the amount of **\$1,072.00** to cover the above fees is enclosed.
- b. ☐ Please charge my Deposit Account No. \_\_\_\_\_ in the amount of \_\_\_\_\_ to cover the above fees. A duplicate copy of this sheet is enclosed.
- c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. **08-2455**. A duplicate copy of this sheet is enclosed.
- d. ☐ Fees are to be charged to a credit card. **WARNING:** Information on this form may become public. **Credit card information should not be included on this form.** Provide credit card information and authorization on PTO-2038.

**NOTE:** Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

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**20741**

PATENT TRADEMARK OFFICE

SIGNATURE

**Christopher J. McDonald**

NAME

**41,533**

REGISTRATION NUMBER

**May 14, 2001**

DATE

09/831355

JC08 Rec'd PCT/PTO 14 MAY 2001

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re the application of:

Michael HOURN; Rodrigo VENTURA;  
John WILLIS and David WINBORNE

ATTN: PCT

Serial No:

Filed :

For : METHOD FOR TREATING PRECIOUS  
METAL BEARING MINERALS

PRELIMINARY AMENDMENT

Commissioner of Patents and Trademarks  
Washington, DC 20231

Sir:

Kindly amend the above-noted application in the  
following manner:

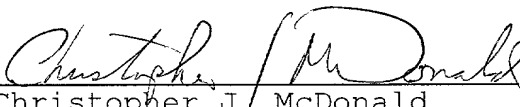
In the Claims:

Please amend the claims as shown on the marked-up set  
of claims and the replacement sheets of claims.

REMARKS

The claims have been changed to place the application  
in conformance with U.S. practice. No new matter has been added.

Respectfully submitted,

by   
Christopher J. McDonald  
Reg. 41,533

May 9, 2001

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Attorney's Docket: A-7419.PAM/cat

CLAIMS:

1. A method of processing a mineral composition comprising a refractory material the method comprising milling the composition to a particle size of  $P_{80}$  of less than 25 microns and leaching said composition with a solution comprising lime and/or limestone in the presence of an oxygen containing gas.

2. The method of claim 1, wherein the refractory material is selected from the group comprising an iron containing sulfide ore, a refinery slime, a carbonaceous ore, a selenide and a telluride.

3. The method of claim 1, wherein the refractory material is selected from the group consisting of pyrite, marcasite, arsenopyrite, troilite, pyrrhotite stibnite, tetrahedrite, argentopyrite, calaverite, altaite, gold bearing selenides, tennantite and pentlandite.

4. The method of claim<sup>3</sup>, where<sup>i</sup>on the refractory material is pyrite or arsenopyrite.

5. The method of claim 2, wherein the material is leached at atmospheric pressure.

6. The method of claim 5 wherein the material is leached in an open tank reactor.

7. The method of claim 2, wherein the material is leached at a temperature of about 50°C up to about the boiling point of the solution.

8. The method of claim 5, wherein the oxygen containing gas is oxygen and the oxygen is introduced into the leaching solution to a level of between about 200 to about 1000kg/tonne of solids in the leaching solution.

CLAIMS:

1. A method of processing a mineral composition comprising a refractory material the method comprising milling the composition to a particle size of  $P_{80}$  of less than 25 microns and leaching said composition with a solution comprising lime and/or limestone in the presence of an oxygen containing gas.

2. The method of claim 1, wherein the refractory material is selected from the group comprising an iron containing sulfide ore, a refinery slime, a carbonaceous ore, a selenide and a telluride.

3. The method of claim 1, wherein the refractory material is selected from the group consisting of pyrite, marcasite, arsenopyrite, troilite, pyrrhotite stibnite, tetrahedrite, argentopyrite, calaverite, altaite, gold bearing selenides, tennantite and pentlandite.

4. (Amended) The method of claim 3, wherein the refractory material is pyrite or arsenopyrite.

5. The method of claim 2, wherein the material is leached at atmospheric pressure.

6. The method of claim 5 wherein the material is leached in an open tank reactor.

7. The method of claim 2, wherein the material is leached at a temperature of about 50°C up to about the boiling point of the solution.

8. The method of claim 5, wherein the oxygen containing gas is oxygen and the oxygen is introduced into the leaching solution to a level of between about 200 to about 1000kg/tonne of solids in the leaching solution.

9. The method of claim 8, wherein the oxygen is introduced into the leaching solution at a flow rate of between about 0.1 to about 0.5vvm.

10. The method of claim 1, wherein the particle size is between about 2 to about 25 microns.

11. The method of claim 1, wherein the particle size is between about 5 to about 15 microns.

12. The method of claim 1, wherein the solution has a pH of the solution is between about 6 to about 12.

13. The method of claim 1, wherein the solution has a pH of between about 6 to about 9.

14. The method of claim 13, wherein the leach solution comprises a mixture of lime and limestone and wt% of limestone in the mixture is between about 40 to about 95%.

15. The method of claim 14, wherein the amount of lime and/or limestone added to the leach solution is between about 100 to about 1200kg/tonne of solids in the solution.

16. The method of claim 15, wherein the amount is about 800kg/tonne.

17. A method of recovering precious metals from a mineral composition comprising a refractory material, the method comprising:

grinding the material to a particle size of 80% passing 25 $\mu$ m or less;

leaching the ground material in the presence of lime and/or limestone and an oxygen containing gas; and

subjecting the leached material to a further leaching step to recover any precious metals.

18. The method of claim 17, wherein the refractory material is a refractory sulfide material bearing gold, silver or platinum.

19. The method of claim 17, wherein the refractory material includes a carbonaceous fraction.

20. The method of claim 17, wherein lime and/or limestone is added to a level of between about 100 to about 1200kg/tonne of solids in the leaching solution and oxygen is introduced to a level of between about 200 to about 1000kg/tonne of solids in the leaching solution.

21. The method of claim 17, wherein the further leaching step uses cyanide as a lixiviant.

22. Gold, silver or platinum recovered by the method of claim 18.

23. A method of recovering gold from a refractory material having a carbonaceous fraction, the method comprising grinding the ore to a particle size of 80% passing 25 micron or less, leaching the ground material with a solution comprising lime and/or limestone at a pH of between about 6 to about 12 in the presence of an oxygen containing gas, subjecting the leached material to a further leaching step in the presence of a cyanide and recovering gold from the cyanide leachate.

24. Gold recovered by the method of claim 23.

## FIELD OF THE INVENTION

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[illegible]



pressure of 2,100 kPa, a pulp density of 20% to 30% solids by mass, acid concentration of 20 - 100 g/L and a retention time of two to three hours. The oxidation must be carried out in an autoclave and requires a source of oxygen.

The capital costs associated with providing the autoclaves and meeting a high oxygen demand are high and may be prohibitive for construction at remote sites, for medium to small scale operations and for low grade ore.

It is possible to leach under less aggressive conditions but in this case, the leaching rates and recovery are too low to be economically viable.

Attempts have been made to reduce the aggressive conditions and to lower the pressures and temperatures required whilst maintaining the economic viability of refractory ore treatment processes. For example, Australian patent application no. 27182/92 describes subjecting an ore concentrate to fine grinding prior to leaching. Fine grinding to an 80% passing size of 15 micron or less enabled the leaching to be carried out under less aggressive conditions at temperatures of 95-110°C and pressures of about 1000 kPa.

Thus, whilst some progress has been made in reducing the operating parameters when using oxygen as the oxidant, the leach must still be carried out under pressure.

US5536480 also describes subjecting an ore to fine grinding prior to acid pressure leaching. In this case, the refractory sulfide ore contains carbonaceous material and the ore is ground to a particle size of 40 microns or less. It was found that in order to obtain an acceptable gold recovery it was necessary to oxidize the material at a minimum temperature of 200°C and to obtain a minimum sulfur oxidation of 96%.

It is also known to oxidatively leach mineral species with ferric ions under acidic conditions. Ferric ion is typically a more effective oxidizing agent than oxygen which means that oxidation with ferric ions can be

carried out under less aggressive conditions. Oxidative leaching at atmospheric pressure using ferric ions is known. A disadvantage of leaching with ferric ions is that the ferric ions are reduced to ferrous ions during the leaching reaction. As leaching solutions are recycled, therefore, ferric ions must be regenerated by oxidizing the reduced ferrous ions.

A further disadvantage with the aforementioned leaching processes for precious metal recovery is that they operate under acidic conditions. One difficulty with leaching under acidic conditions is that ores which contain an organic carbon fraction, known as carbonaceous ores cannot readily be processed to obtain acceptable levels of precious metal recovery. Precious metals such as gold are typically recovered from a leach solution by a cyanide leach stage. The reduction in recovery is due to absorption of the precious metals by the organics, during the cyanide stage. The absorbed metals cannot be recovered by cyanidation without a further pre-treatment stage designed to destroy the carbonaceous matter.

In order to address the problem of organic carbon, carbonaceous refractory ores are commonly treated by roasting to convert carbonaceous matter in the ores to gaseous carbon dioxide, or by treatment with a strong chemical oxidant, such as chlorine, to oxidize the carbonaceous matter. Both methods are expensive and are not economically viable for treatment of low grade materials.

Still further, as leaching steps such as cyanidation require alkaline conditions, the acid must be removed prior to cyanidation.

The problem of removing acid prior to cyanidation may be overcome by leaching under alkaline conditions. Leaching of nickel and cobalt under alkaline conditions using ammonia/ammonium salts is well known. However, a major disadvantage of alkaline leaching is that when iron containing ores such as pyrite are oxidised, the iron which is leached precipitates as a

passive iron oxide layer on the mineral particle. This layer inhibits further oxidation with the result being that the extent of leaching under alkaline conditions is less than under acidic conditions. This translates to a lower recovery of precious metals.

Still further, alkaline leaching of refractory materials requires elevated pressure and temperatures and an oxidant for the leaching to occur. However, even under aggressive alkaline conditions, recovery of precious metals is often less than that for acid leaching. Further, base metals such as copper and zinc are insoluble at high pH. Thus, alkaline leaching is unsuitable for leaching ores or concentrates where recovery of base metals from base metal sulphides such as chalcocite, sphalerite or chalcopyrite is required. For these reasons, commercial and academic interest has been directed towards acid leaching.

Most of the literature relating to alkaline leaching is directed towards the use of water soluble alkalis such as sodium or potassium hydroxide and ammonia. A disadvantage with these reagents is that iron is precipitated primarily as jarosite. Jarosite inhibits gold recovery and is also an environmentally unacceptable residue. Also, hydroxide reagents and in particular sodium hydroxide are prohibitively expensive.

The use of cheaper alkalis such as lime has been proposed. However, to date, leaching of iron sulphide materials with lime has been unsuccessful in that leaching is incomplete and subsequent precious metal recovery is low. For example, an earlier study of alkaline oxidation of pyrite for gold recovery using lime achieved only 30 to 40% gold recovery which offered little improvement over direct cyanidation of the pyrite. This is believed to be due to passivation of the mineral by precipitation of a gypsum/iron oxide layer.

Limestone is another alkali which is relatively cheap. Limestone is typically used in the neutralization of acidic leachates. However, limestone is considered to

be insufficiently reactive and/or soluble in alkaline systems to be able to be used for alkaline leaching.

From an economic point of view, it would be desirable to be able to leach refractory materials for precious metal recovery under mild alkaline conditions and using reagents other than the expensive hydroxides.

As mentioned above, it is known that the oxidation rate under acidic conditions can be increased by fine grinding to increase the surface area of the mineral particles. Such an increase may be predicted given that there is a larger surface area exposed to the oxidizing agents. However in the alkaline system, this effect is substantially reduced in view of the formation of the passive iron oxide layer on the particles. The rate determining factors in the alkaline systems are believed to relate to the formation of the passivating iron oxide layer and diffusion of oxygen through the layer. Thus, workers in the field have concentrated on increasing the extent of alkaline leaching by using strong, soluble alkalis, by modifying the leaching conditions so as to minimise formation of the passive layer and/or influence the diffusion rate through the layer.

One study suggests leaching at higher temperatures or at relatively concentrated solutions of reagents. The reason for this is to rapidly produce a passive layer which is unstable and subject to cracking.

It is believed that at lower temperatures, the layers grow more slowly and are more stable. Another suggestion has been to use additives which may react to dissolve the layer or to make the layer more permeable.

However, to date no method has been proposed which is able to economically leach iron containing refractory ores and concentrates under alkaline conditions and which also enables good recovery of precious metals from the ore or concentrates.

The present invention is based on the surprising and unexpected discovery that leaching

refractory sulfide and/or carbonaceous materials under alkaline conditions can be successfully achieved by careful selection of the particle size of the material to be leached. Even more surprising it has been discovered that not only can the leaching efficiency be improved but that leaching can also be successfully conducted under relatively mild conditions. Still further, it has also been discovered that the activity of any carbonaceous matter in the feed material can also be substantially reduced as part of the oxidation process, to a point where it will not compete with the activated carbon added to commercial gold/silver recovery processes, where cyanide is used to leach the precious metals.

According to a first broad form of the invention there is provided a method of processing a mineral composition comprising a refractory material the method comprising milling the composition to a particle size of  $P_{80}$  of less than 25 microns and leaching said composition with a solution comprising lime and/or limestone in the presence of an oxygen containing gas.

In the present specification and claims the term "refractory material" includes refractory sulfides such as pyrite in which precious metals are encapsulated, ores which contain carbonaceous material and telluride or selenide materials. In the present specification and claims, the use of the term "ore" Includes not only ore per se but also includes concentrates, slimes, tailings, spoil and waste materials which may have a recoverable amount of precious metal values. Carbonaceous material refers to materials having an organic carbon fraction which may include graphite, bituminous or partly bituminous material.

The method of the present invention is particularly applicable to a composition including an iron containing refractory sulphide, selenide or telluride material or a mixture of such materials, with or without carbonaceous material present. Examples of such materials include pyrite, marcasite, arsenopyrite

and arsenic bearing pyrite, troilite and pyrrhotite.

According to a further broad form of the invention there is provided a method of processing a mineral composition comprising an iron containing refractory material the method comprising milling the composition to a particle size of  $P_{80}$  of less than 25 microns and leaching said composition with a solution comprising lime and/or limestone in the presence of an oxygen containing gas.

Other non-iron or minor iron containing materials may also be present in the composition, examples of which include stibnite, tetrahedrite, argentopyrite, calaverite, altaite, gold bearing selenides, tennantite and pentlandite. The method of the present invention is also applicable to a composition including carbonaceous matter, where the carbonaceous matter would otherwise interfere with the precious metals recovery process. Suitably the composition would not include economic amounts of base metal sulphides containing copper or zinc. Generally a composition which includes appreciable amounts of these base metals would not be treated directly by the method of the present invention. The reason for this is purely economical as copper and zinc precipitate under alkaline conditions and thus cannot be recovered easily. Under acidic leaching conditions, copper and zinc are solubilized and can be recovered by conventional SX/EW techniques. Use of flotation or other separation technologies to produce a base metal concentrate and a separate refractory iron sulphide concentrate would be obvious to one skilled in the art as a way of treating these materials.

The method of the present invention is in particular directed towards the treatment of refractory materials containing precious metals such as gold, silver and platinum. Suitably, the refractory materials are in the form of flotation concentrates although the method is suitable for ores if the economics are favourable.

In the method of the present invention, the

composition is finely ground to a particle size of 80 % by mass passing less than 25 $\mu$ m. A typical particle size range is between 80 % passing 2-25 $\mu$ m and preferably between about 80 % passing 5-15 $\mu$ m.

5           A preferred apparatus for producing the finely ground material is a stirred ball mill. However, it will be appreciated that several other suitable types of comminution apparatus may also be used.

10           The leaching is carried out using lime and/or limestone as the alkali reagent. Lime, limestone or a mixture thereof may be used. Preferably a mixture in the range of 40-95% limestone is used. The lime and/or limestone is added in an amount such that the pH of the system is between about 6-12 and preferably about 6-9.

15           Typically about 100 to about 1200kg, of lime and/or limestone is added per tonne of solids. The amount of lime and/or limestone which will need to added to maintain a desired pH will generally vary according to the amount (if any) of sulphuric acid produced by  
20 sulphide oxidation. Generally about 800kg of lime and/or limestone would be added.

          The present inventors have also surprisingly discovered that not only can the overall amount of leaching be increased but that such an increase can also  
25 be obtained by leaching under milder conditions than has hitherto been possible. Suitably the method of the present invention can be carried out at ambient pressure. This avoids the use of expensive pressure reactors and autoclave equipment. The preferred operating temperature  
30 is between about 50°C up to the boiling point of the mixture. Typically the maximum temperature is about 95°C.

          The leaching reaction can thus be carried out in open tank reactors. Excess heat is removed by  
35 evaporation of the solution. This avoids the need for costly heat exchangers. Heat can be easily introduced by known methods such as the injection of steam.

          The leaching reaction is carried out in the

presence of an oxygen containing gas. When the reaction is carried out in an open tank reactor the gas is typically introduced by sparging. The gas may be oxygen, air or oxygen enriched air. The gas flow is dependent upon the amount of oxygen required to sustain the leaching reaction and is typically about 0.01 - 0.5 vvm (vessel volumes per minute). The oxygen consumption of the process is typically between about 200 - 1000kg of oxygen per tonne of solids.

After the composition has been leached, the mixture can be further treated by known methods to recover precious metals, principally by cyanide leaching.

Cyanidation occurs under alkaline conditions. As the leach solution is already alkaline, the solution may be subjected directly to cyanide leaching. If desired the slurry may be thickened prior to cyanide leaching. A further advantage of the current invention is that the type of compounds that form in the alkaline leaching system are not reactive toward cyanide, and will not consume high levels of cyanide in the gold recovery stage. Compounds formed under acidic leach conditions often consume significant amounts of cyanide in the gold recovery stage, increasing the process costs.

According to a further broad form of the invention there is provided a method of recovering precious metals from a mineral composition comprising a refractory material, the method comprising;

grinding the material to a particle size of 80 % passing 25 $\mu$ m or less;

leaching the ground material in the presence of lime and/or limestone and an oxygen containing gas; and

subjecting the leached material to a further leaching step to recover any precious metals.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a flow diagram of a preferred method of the present invention and

Figure 2 is a flow diagram of a further preferred method of the present invention.



## BEST MODE

Referring now to Figure 1, the ore or concentrate 10 is slurried to about 50% solids and fed into a stirred ball mill 11 and milled to a particle size of 80% passing 25  $\mu\text{m}$  or less. The milled material is then fed into a non-pressurized reactor 12. Oxygen or air 14 is introduced into the reactor 12 and leaching is carried out at atmospheric pressure at a temperature of between about 50 to about 95°C. Lime and/or limestone 13 are added to the reactor 12 to control the pH to between about 6 and about 12. The leached material is then subjected to metals recovery 14.

Figure 2 illustrates a further flow diagram. This diagram is similar to that of Figure 1 and the same reference numerals have been used to identify the same steps or reagents. The process illustrated in this diagram further includes a thickener 15 which thickens the slurry following the leaching stage in reactor 12 prior to metals recovery 14. The excess solution 16 is returned to the mill 11 for reuse.

The present invention will now be described in relation to the following examples.

**Example No. 1. Leach of a pyrite containing ore, with a 50:50 blend of limestone and lime, oxygen used as the oxidant.**

**Alkali Leach**

The ore sample was slurried in tap water at 50% solids and milled in a laboratory rod mill for a period of 20 minutes to achieve a slurry ground to 80% passing 106 microns. The slurry was then thickened to 60% solids, and milled in a Netzsch horizontally stirred laboratory scale bead mill. The media used was 0.8 - 1.1mm steel balls. Slurry was passed through the mill sequentially to achieve the required grind size of 80% passing 14 microns. The final particle size distribution was determined by lasersizer.

The leaching test was carried out in a cylindrical stainless steel reactor fitted with four baffles located equidistant around the edges of the vessel. A 1,000 gram sample of the ground solids were  
5 added to the leach vessel along with 20 litres of tap water. The slurry was agitated by a 6 bladed Rushton style impeller. Oxygen was introduced to the reactor by an air spear, which terminated below the impeller. The oxygen flow was controlled by a rotameter at 2,000 cubic  
10 centimetres per minute (0.1 vvm).

The pH in the vessel was maintained by an automated pH analyser and controller, to a set point of 10. A combination glass pH probe was immersed in the slurry through a port in the lid of the reaction vessel.  
15 A limestone/lime slurry was continuously circulated from a stirred beaker through a ring main using a peristaltic pump. The limestone/lime slurry consisted of 50% lime and 50% limestone, at a slurry density of 35% w/w solids.

A solenoid valve was attached to the ring main, with an outlet line directed to the reaction vessel. The  
20 controller actuated the solenoid when the pH in the vessel drifted below the set point.

The vessel was heated by steam injection into a jacket surrounding the vessel. The steam injection was  
25 regulated by a solenoid valve, which was controlled by an automatic temperature controller. The vessel temperature was controlled to 85°C. The outside of the vessel was wrapped in insulated foam to prevent heat loss.

On completion of the test, the slurry was  
30 filtered and the filter cake dried and weighed. A sample of the filter cake was analysed for residual sulphide by acid evolution. A further sample of the cake was sent for XRD analysis to confirm the sulphide analysis. The dry weight of the filter cake was 1150 grams. The results of  
35 the sulphide oxidation step are listed below in Table 1.

Table 1

Degree of sulphide Oxidation Achieved for Whole Ore Leach

Sample	Weight - grams	% sulphide sulphur	% Pyrite by sulphide evolution	% Pyrite by XRD	% sulphide oxidation
Feed	1000	8.2	15.2	15.8	90.1
Leach	1150	0.7	1.30	1.6	
Residue					

Example No. 2. Comparative leaching of a pyrite  
5 containing concentrate, with a 80:20 blend of limestone  
and lime and acid, oxygen used as the oxidant.

#### Alkali Leach

10 The concentrate sample was milled according to  
the procedure outlined in example 1, to achieve the  
required grind size of 80% passing 9.6 microns.

The leaching test was carried out in a 10 litre  
cylindrical stainless steel reactor fitted with four  
baffles located equidistant around the edges of the  
15 vessel. A 2000 gram sample of the ground solids was  
added to the leach vessel along with 10 L of tap water.

The slurry was agitated by a 6 bladed Rushton  
style impeller. Oxygen was introduced to the reactor by  
an air spear, which terminated below the impeller. The  
20 oxygen flow was controlled by a rotameter at 800 cubic  
centimetres per minute (0.08 vvm).

A single addition of 2100 grams of limestone  
and 380 grams of hydrated lime was made to the vessel at  
the start of the test.

25 The vessel was placed on a thermocouple  
controlled hotplate, with the thermocouple immersed in  
the slurry through a stainless steel sheath. The vessel  
was maintained at the required temperature by the  
thermocouple controller. The temperature was controlled  
30 to 82°C.

On completion of the test, the slurry was  
filtered and the filter cake dried and weighed. A sample

of the filter cake was analysed for residual sulphide. The dry weight of the filter cake was 4230 grams.

#### Acid Leach

5           The concentrate sample was milled according to the procedure outlined in Example 1, to produce a final product at 80 % passing 9.6 microns. The leaching test was carried out in a 10 litre cylindrical stainless steel reactor fitted with four baffles located equidistant  
10 around the edges of the vessel. A 1400 gram sample of the ground solids was added to the leach vessel along with 10 L of tap water. The slurry was agitated by a 6 bladed Rushton style impeller. Oxygen was introduced to the reactor by an air spear, which terminated below the  
15 impeller. The oxygen flow was controlled by a rotameter at 800 cubic centimetres per minute (0.08 vvm).

A single addition of 100 grams of sulphuric acid and 250 grams of ferric sulphate hexahydrate was made to the vessel at the start of the test. The  
20 vessel temperature was controlled to 82°C. On completion of the test, the slurry was filtered and the filter cake dried and weighed. A sample of the filter cake was analysed for residual sulphide. The dry weight of the filter cake was 1812 grams.

25           The results of the sulphide oxidation steps are listed below in Table 2.

**Table 2**  
**Degree of Sulphide Oxidation Achieved for Pyrite**  
**Concentrate**

Sample	Weight - grams	% sulphide sulphur	% sulphide oxidation
Alkaline Leach			
Feed	2000	42	88.5
Leach	4230	2.3	
Residue			
Acid Leach			
Feed	1400	42	74.7
Leach	1812	8.2	
Residue			

### Cyanide Leach

A sample of each leach residue and the un-oxidised feed material were slurried in tap water at a slurry density of 40% w/w solids in a baffled glass reactor. The volume of the reactor was 3 litres. The slurry was agitated by a 6 bladed Rushton downdraft style impeller. Air was introduced to the reactor by the action of the downdraft impeller.

The pH of the slurry was adjusted and maintained at pH 10 using lime, and sodium cyanide was added to the slurry to maintain a free cyanide level of 500ppm. The test was carried out for a period of 24 hours. On completion of the test, the slurry was filtered and the filter cake washed with deionised water.

The filtrate and washings were combined for analysis. The solid filter cake was dried in an oven and analysed to determine the precious metals recovery. The results of the cyanide leach are listed in Table 3.

Table 3.

## Cyanide Leach Results for Oxidised Pyrite Concentrate

Sample	% pyrite oxidised	Gold Extraction %	Sodium Cyanide consumed - kg/tonne
Feed prior to oxidation	NA	26.6	4.5
Acid Oxidized Residue	74.7	72	11.7
Alkaline Oxidised Residue	88.5	94	3.6

5           The precious metals extractions were determined  
by head and tails fire assay.

10       **Example No. 3. Leach of a pyrite containing concentrate,**  
with a 80:20 blend of limestone and lime, air used as the  
oxidant.

**Alkali Leach**

15       The concentrate sample was milled according to  
the procedure outlined in example 1, to achieve the  
required grind size of 80% passing 12 microns. The final  
particle size distribution was determined by lasersizer.

20       The leaching test was carried out in a  
cylindrical stainless steel reactor fitted with four  
baffles located equidistant around the edges of the  
vessel. A 200 gram sample of the ground solids were  
added to the leach vessel along with 2,000mL of tap  
water. The live volume of the reactor was 2.5 litres.

The vessel was fitted with a stainless steel lid to prevent evaporative loss from the reactor. The slurry was agitated by a 6 bladed Rushton style impeller. Air was introduced to the reactor by an air spear, which  
5 terminated below the impeller. The air flow was controlled by a rotameter at 200 cubic centimetres per minute (0.1 vvm).

The pH in the vessel was maintained using the same system described in Example 1, with the exception  
10 that the limestone/lime slurry consisted of 20% lime and 80% limestone, at a slurry density of 35% w/w solids.

The vessel was placed on a thermocouple controlled hotplate, with the thermocouple immersed in the slurry through a stainless steel sheath. The vessel  
15 was maintained at 85°C by the thermocouple controller.

On completion of the test, the slurry was filtered and the filter cake dried and weighed. A sample of the filter cake was analysed for residual sulphide by acid evolution. The dry weight of the filter cake was 469  
20 grams. The results of the sulphide oxidation step are listed below in Table 4.

Table 4

Degree of Sulphide Oxidation Achieved for Pyrite  
25 Concentrate when oxidised with Air as the Oxidant

Sample	Weight - grams	% sulphide sulphur	% Pyrite by acid evolution	% sulphide oxidation
Feed	200	33.2	61.73	90.1
Leach	469	1.4	2.6	
Residue				

**Example No. 4. Leach of an arsenopyrite containing concentrate, with a 50:50 blend of limestone and lime, oxygen used as the oxidant.**

#### 5 Alkali Leach

An arsenopyrite concentrate, grading 10.73 % arsenic and 32 % sulphur was tested for gold recovery. The concentrate sample was milled according to the procedure outlined in example 1, to achieve the required grind size of 80% passing 14 microns. The final particle size distribution was determined by lasersizer.

The leaching test was carried out according to the procedure outlined in Example 1. A 1500 gram sample of the ground solids were added to the leach vessel. The pH in the vessel was maintained according to the method outlined in Example 1.

On completion of the test, the slurry was filtered and the filter cake dried and weighed. A sample of the filter cake was analysed for residual sulphide by acid evolution. A further sample of the cake was sent for XRD analysis to confirm the sulphide analysis. The dry weight of the filter cake was 2965 grams. The results of the sulphide oxidation step are listed below in Table 5.

25

**Table 5**

**Degree of Sulphide Oxidation Achieved for Arsenopyrite Concentrate**

Sample	Weight grams	% sulphide sulphur	% Arsenopyrite by XRD	% Pyrite by XRD	% sulphide oxidation
Feed	1500	20.1	30.6	29.5	93.1
Leach Residue	2965	0.7	<1	1.4	

The presence of arsenic as ferric arsenate in the leach residue was confirmed by XRD

30



### Cyanide Leach

The cyanide leach procedure was the same as outlined for Example 2. The results of the cyanide leach are listed in Table 6.

5

Table 6.  
Cyanide Leach Results for Oxidised Pyrite and  
Arsenopyrite Concentrate

Sample	% pyrite oxidised	% arsenopyrite oxidised	Gold Extraction %
Feed	NA	NA	35.8
Oxidised Residue	90.6	>95	88.05

10

The precious metals extractions were determined by head and tails fire assay.

### Example No. 5. The Effects of Varying Alkali Mixtures on the Extent of Sulphide Oxidation

15

#### Alkali Leach

Approximately 5 kg of pyrite concentrate was milled according to the procedure outlined in example 1, to achieve the required grind size of 80% passing 8.6 microns. The final particle size distribution was determined by lasersizer. On completion of the grind, the ground slurry was split into 500 gram subsamples for the leaching test, and each subsample filtered separately. All subsamples were stored frozen to prevent oxidation of the pyrite.

20

25

A 200 gram sample was split out of each subsample for the leaching testwork. The leaching test was carried out according to the procedure outlined in Example 3. A 200 gram sample of the ground solids was used for the test. The pH in the vessel was maintained using the same system described in Example 1.

30

The limestone/lime slurry composition was varied for the three leach tests according to the following ratios: 100% lime, 50% lime 50% limestone, 10% lime 90% limestone. The vessel temperature was controlled to 85°C. On completion of the test, the slurry was filtered and the filter cake dried and weighed. A sample of the cake was sent for XRD analysis to determine the extent of pyrite leaching. The dry weight of the filter cake was recorded.

The results are listed in Table 7, below

**Table 7.**  
**The Effects Of Varying Lime/Limestone Blends On The**  
**Extent Of Pyrite Oxidation From Pyrite Flotation**  
**Concentrate**

Alkaline Leach Tests			
PH	10	10	10
Alkali mixture	100% lime	50% lime 50% limestone	10% lime 90% limestone
Grind Size - 80% passing	8.6	8.6	8.6
% solids	10.0	10.0	10.0
Feed data:			
Dry solids wt g	200	201.0	201.1
% pyrite in head	65.0	65.0	65.0
Residue data:			
% pyrite in residue	3.2	3.0	1.0
% pyrite oxidised from XRD	95.47	90.1	94.3

**Example No. 6. The Effects of Varying Grind Size on the Extent of Sulphide Oxidation.**

**Alkali Leach**

5 Approximately 4 kg of pyrite concentrate sample was milled according to the procedure outlined in example 1, to produce 6 individual samples at the required grind sizes of 80% passing:

10	Pass1	51.9 microns
	Pass2	32.04 microns
	Pass3	17.94 microns
	Pass4	13.64 microns
	Pass5	11.71 microns
15	Pass6	8.6 microns

On completion of the grind, the ground slurry was split into 500 gram subsamples for the leaching test, and each subsample filtered separately. All subsamples  
20 were stored frozen to prevent oxidation of the pyrite.

A 200 gram sample was split out of each subsample for the leaching testwork. The leaching tests were carried out according to the procedure outlined in Example 3. The pH in the vessel was maintained using the  
25 system described in Example 1. The vessel temperature was controlled to 80°C. On completion of the test, the slurry was filtered and the filter cake dried and weighed. A sample of the cake was sent for XRD analysis to determine the extent of pyrite leaching. The dry weight of the  
30 filter cake was recorded.

The results are listed in Table 8, below

Table 8.

The Effects of Varying Grind Size on the Extent of Pyrite Oxidation from Pyrite Flotation Concentrate

5

Alkaline Leach Tests						
pH	10	10	10	10	10	10
Alkali mixture	100%	100%	100%	100%	100%	100%
	lime	lime	lime	lime	lime	lime
Grind Size - 80% passing	51.9	32.04	17.94	13.64	11.71	8.6
% solids	10.0	10.0	10.0	10.0	10	10.0
Feed data:						
Dry solids wt g	200.0	200	200	200	200	200
% pyrite in head	65.0	65.0	65.0	65.0	65.0	65.0
Residue data:						
% pyrite in residue	16.7	9.7	7	2.7	3.7	3.2
% pyrite oxidised from XRD	73.9	80.1	91.06	96.81	95.54	95.47
% additional gold recovery	0	4	8	12	14	14

Example No. 7. Leach of a pyrite containing concentrate, with a 50:50 blend of limestone and lime with oxygen used as the oxidant, at varying pH.

10

#### Alkali Leach

The concentrate sample was milled according to the procedure outlined in example 1, to achieve the required grind size of 80% passing 12 microns. All leaching tests were carried out according to the procedure outlined in Example 2.

15

The pH in the vessel was maintained by an automated pH analyser and controller. In all, three tests were carried out, each at a different pH. The pH control points used were

5

Test 1	pH 8
Test 2	pH 9
Test 3	pH 10

The limestone/lime slurry consisted of 50% lime and 50% limestone, at a slurry density of 35% w/w solids. The vessel temperature was controlled to 85°C.

10

On completion of the test, each slurry was filtered and the filter cake dried and weighed.

**Table 9.**

**Degree of oxidation of Pyrite at Varying pH**

15

Control pH	Not oxidised	Oxidised at pH 8	Oxidised at pH 9	Oxidised at pH 10
% sulphur oxidation	<10	90	94	94

#### Cyanide Leach

The cyanide leach procedure was the same as outlined for Example 2. The results of the cyanide leach are listed in Table 10.

20

**Table 10.**

**Gold Recoveries from Oxidised Pyrite concentrate at Varying pH**

25

Control pH	Not oxidised	Oxidised at pH 8	Oxidised at pH 9	Oxidised at pH 10
Au recovery - %	26.6	92.2	96.1	97.2

The precious metals extractions were determined by head and tails fire assay.

**Example No. 8. Leach of a pyrite concentrate containing high levels of carbonaceous matter.**

**5 Alkali Leach**

A 1200 gram sample of concentrate containing approximately 18 % w/w pyrite and 21.8 % w/w organic carbon was produced by flotation of a carbonaceous ore sample with diesel. The concentrate sample was milled according to the procedure outlined in Example 1, to achieve the required grind size of 80% passing 9 microns

The leach test was carried out according to the procedure outlined in Example 3, using a 120 gram sample of the ground concentrate. A single addition of 40 grams of hydrated lime and 160 grams of limestone was made to the leach at the start of the leach test, and the slurry allowed to react for 24 hours. The vessel temperature was controlled to 85°C. On completion of the test, the slurry was filtered and the filter cake dried and weighed.

**Gold Adsorption Tests**

A 20 ppm gold in cyanide solution was made up in de-ionised water. The free cyanide level in the solution was 500 ppm.

A 20 gram sample of the oxidised concentrate was added to 500 mL of the gold cyanide solution, and the solution agitated by magnetic stirrer. The solution was sampled regularly over a period of 75 minutes. A 20 gram sample of the ground concentrate, which had not been subjected to oxidative leaching, was also added to 500 mL of gold cyanide solution, and the solution agitated by magnetic stirrer. The solution was also sampled regularly over a period of 70 minutes.

All solution samples were analysed for gold by DIBK extract/AAS. The results of the gold adsorption tests are listed below in Table 11

**Table 11**  
**Effect of Oxidative Alkaline Leaching on the Activity of**  
**Carbonaceous Material.**

Time (min)	Gold in Solution (ppm)	
	Alkaline leach residue	Ground Concentrate
0	17.4	15.9
1	15.2	11.4
5	15.2	10.2
10	14.9	9.65
20	14.8	9.20
50	14.6	9.10
70	15.4	-
75	-	9.20

**Example No. 9. Leach of Electrolytic Copper Refinery Slimes Containing Gold And Silver Selenides and Tellurides.**

#### **Alkali Leach**

A 1500 gram sample of Electrolytic Copper Refinery Slimes, assaying 16.2% Cu, 8400g/t Au, 7.8% Ag, 0.24% Te, 2.45% Se, was milled according to the procedure outlined in Example 1, to 80% passing 9 microns.

Copper Refinery Slimes contain gold and silver in the form of telluride and selenides. Typical constituents of the slimes include phases with the composition  $(\text{Cu,Ag})_2\text{Se}$  and  $(\text{Cu,Ag})_2\text{Te}$ . These phases do not leach in conventional cyanide leach circuits.

The leach test was carried out according to the procedure outlined in Example 2. A 570 gram sample of the ground slimes was added to the leach vessel along with 8 L of tap water. A single addition of 380 grams of hydrated lime and 1520 grams of limestone was made to the leach at the start of the leach test, and the slurry allowed to react for 24 hours. The vessel temperature was controlled to 85°C. On completion of the test, the slurry was filtered and the filter cake dried and weighed.

### Cyanide leaching Tests

Two cyanide leaching tests were carried out. The cyanide leach procedure was the same as outlined for Example 2. The results of the cyanide leach are listed in Table 12.

Table 12.

#### Gold Recoveries from Copper Refinery Slimes

	Alkaline Leached slimes	Non Alkaline Leached Slimes
Gold Extraction	92	46.5

Gold recovery from the alkaline leached slimes was significantly higher than for the fresh slimes, indicating that the gold bearing selenide and telluride phases present in the slimes were broken down in the alkaline leach.

#### Example No. 10. Leach of a Stibnite ( $\text{Sb}_2\text{S}_3$ ) concentrate.

##### Alkali Leach

A 250 gram sample of concentrate containing 40 % w/w  $\text{Sb}_2\text{S}_3$  and 60 % w/w siliceous gangue was slurried in tap water at 60% solids, and milled in an ECC vertically stirred laboratory mill. Slurry was milled to achieve the required grind size of 80% passing 12 microns. The final particle size distribution was determined by lasersizer.

The leach test was carried out according to the procedure outlined in Example 3. A 200 gram sample of the ground concentrate was used in the test.

A single addition of 30 grams of hydrated lime and 120 grams of limestone was made to the leach at the start of the leach test, and the slurry allowed to react for 24 hours. The vessel temperature was controlled to 85°C. On completion of the test, the slurry was filtered and the filter cake dried and weighed. The results of the oxidation test are listed in Table 12, below.



Table 12

Oxidation of finely ground Stibnite ( $\text{Sb}_2\text{S}_3$ ) with  
lime/limestone under alkaline conditions.

Sample	Weight	% sulphide	% sulphide oxidation
Feed concentrate	200	9.43	87
Oxidised Residue	326.9	0.76	

5

It can be seen that by grinding to  $25\mu\text{m}$  or less and leaching under alkaline conditions with lime and/or limestone that 90% or more of the refractory components of the feed can be oxidised. A high level of oxidation typically translates to high precious metal recovery. However, for some ores or concentrates a high level of gold recovery can be achieved with comparatively low levels of sulfide oxidation. When processing such ores or concentrates according to the invention, typically only, as much sulfur as required would be oxidized to obtain a desired level of gold recovery.

The high degree of oxidation of the minerals which can be achieved as a result of the present invention sets it apart from other processes where air and lime are added to sulphide slurries prior to cyanide leaching. In these other processes, the addition of air to a level of about 10-20 kg/tonne solids and lime, to a level of only 5 - 20 kg/tonne of solids, is designed to remove soluble ions from the solution phase of the slurry that may impact adversely on the cyanide leach stage. The current invention, on the other hand, substantially breaks down the solid phase of the slurry, liberating encapsulated gold for recovery through cyanide leaching, with much higher consumption of lime and or limestone.

The methods of the present invention offer a number of advantages over existing methods. First, iron containing refractory ores such as pyrite and

arsenopyrite can be oxidised to high levels under alkaline conditions using lime and/or limestone as the alkali source. Selenides and Tellurides present in these feeds are broken down in the process and the activity of any carbonaceous matter in the feed is substantially reduced. Lime and limestone offer significant economic advantages over the known agents such as caustic. For example, current costs for caustic are about AUD\$440 per tonne, ammonium based salts, about AUD\$450, lime AUD\$100-200 and limestone AUD\$15-41/tonne.

Further, the leach need not be pressurized, which translates to significant capital and operating costs. Still further, costs can be reduced as the leach can be carried out using air instead of oxygen. This means that oxygen need not be purchased or produced. Further economic benefits can be realized as the cyanide consumption of residues leached by the present method is less than that for residues leached under acidic conditions.

The leach residence times for the preferred methods of the present invention are typically about 12 - 30 hours. These residence times compare favorably to oxidative leaching under acidic conditions and are in fact superior to atmospheric ferric leaching.

Further, leaching under the condition of the present invention inactivates any carbonaceous material which allows precious metals to be recovered by cyanide lixiviation.

During the leaching iron precipitates as goethite and hematite, rather than Jarosite as occurs under acidic conditions. Jarosite inhibits subsequent precious metal recovery by making the residue difficult to settle and filter. Further, Jarosite is not an environmentally acceptable residue.

Gypsum is also formed during the leach and precipitates. An advantage of gypsum is that it enhances the filterability of the residue. Gypsum is not formed when using conventional alkali reagents.

Another advantage over the use of the water soluble alkali reagents is that in the present invention arsenic present in the refractory material is precipitated as ferric arsenate. The level of arsenic in the leach liquor is typically below detectable limits. When using conventional soluble alkalis arsenic is present in the leach liquor.

The process of the present invention has enabled recovery of precious metals from sulfide and/or carbonaceous ores under (1) conditions of temperature and pressure which were previously believed to be insufficiently reactive for such ore materials, (2) the use of alkaline reagents which were also previously believed to be insufficiently reactive and/or soluble and (3) under pH conditions which were previously believed to lead to passivation and incomplete oxidation of the ore particle.

In the present specification and claims, the term "comprise" and variations such as "comprises" and "comprising" or the term "include" or variations thereof will be understood to imply the inclusion of a stated element or integer or a group of integers or elements, but not the exclusion of any other element or integer or group of elements or integers.

## CLAIMS

1. A method of processing a mineral composition comprising a refractory material the method comprising milling the composition to a particle size of  $P_{80}$  of less  
5 than 25 microns and leaching said composition with a solution comprising lime and/or limestone in the presence of an oxygen containing gas.
2. The method of claim 1, wherein the refractory material is selected from the group comprising an iron  
10 containing sulfide ore, a refinery slime, a carbonaceous ore, a selenide and a telluride.
3. The method of claim 1, wherein the refractory material is selected from the group consisting of pyrite, marcasite, arsenopyrite, troilite, pyrrhotite stibnite,  
15 tetrahedrite, argentopyrite, calaverite, altaite, gold bearing selenides, tennantite and pentlandite.
4. The method of claim, whereon the refractory material is pyrite or arsenopyrite.
5. The method of claim 2, wherein the material is  
20 leached at atmospheric pressure.
6. The method of claim 5 wherein the material is leached in an open tank reactor.
7. The method of claim 2, wherein the material is leached at a temperature of about 50°C up to about the  
25 boiling point of the solution.
8. The method of claim 5, wherein the oxygen containing gas is oxygen and the oxygen is introduced into the leaching solution to a level of between about 200 to about 1000kg/tonne of solids in the leaching  
30 solution.
9. The method of claim 8, wherein the oxygen is introduced into the leaching solution at a flow rate of between about 0.1 to about 0.5vvm.
10. The method of claim 1, wherein the particle  
35 size is between about 2 to about 25 microns.
11. The method of claim 1, wherein the particle size is between about 5 to about 15 microns.
12. The method of claim 1, wherein the solution has a pH

of the solution is between about 6 to about 12.

13. The method of claim 1, wherein the solution has a pH of between about 6 to about 9.

14. The method of claim 13, wherein the leach  
5 solution comprises a mixture of lime and limestone and wt% of limestone in the mixture is between about 40 to about 95%.

15. The method of claim 14, wherein the amount of lime and/or limestone added to the leach solution is  
10 between about 100 to about 1200kg/tonne of solids in the solution.

16. The method of claim 15, wherein the amount is about 800kg/tonne.

17. A method of recovering precious metals from a  
15 mineral composition comprising a refractory material, the method comprising;

grinding the material to a particle size of 80% passing 25 $\mu$ m or less;

leaching the ground material in the presence of  
20 lime and/or limestone and an oxygen containing gas; and  
subjecting the leached material to a further leaching step to recover any precious metals.

18. The method of claim 17, wherein the refractory material is a refractory sulfide material bearing gold,  
25 silver or platinum.

19. The method of claim 17, wherein the refractory material includes a carbonaceous fraction.

20. The method of claim 17, wherein lime and/or limestone is added to a level of between about 100 to  
30 about 1200kg/tonne of solids ion the leaching solution and oxygen is introduced to a level of between about 200 to about 1000kg/tonne of solids in the leaching solution.

21. The method of claim 17, wherein the further leaching step uses cyanide as a lixiviant.

35 22. Gold, silver or platinum recovered by the method of claim 18.

23. A method of recovering gold from a refractory material having a carbonaceous fraction, the method

- comprising grinding the ore to a particle size of 80% passing 25 micron or less, leaching the ground material with a solution comprising lime and/or limestone at a pH of between about 6 to about 12 in the presence of an oxygen containing gas, subjecting the leached material to a further leaching step in the presence of a cyanide and recovering gold from the cyanide leachate.
- 5
24. Gold recovered by the method of claim 23.

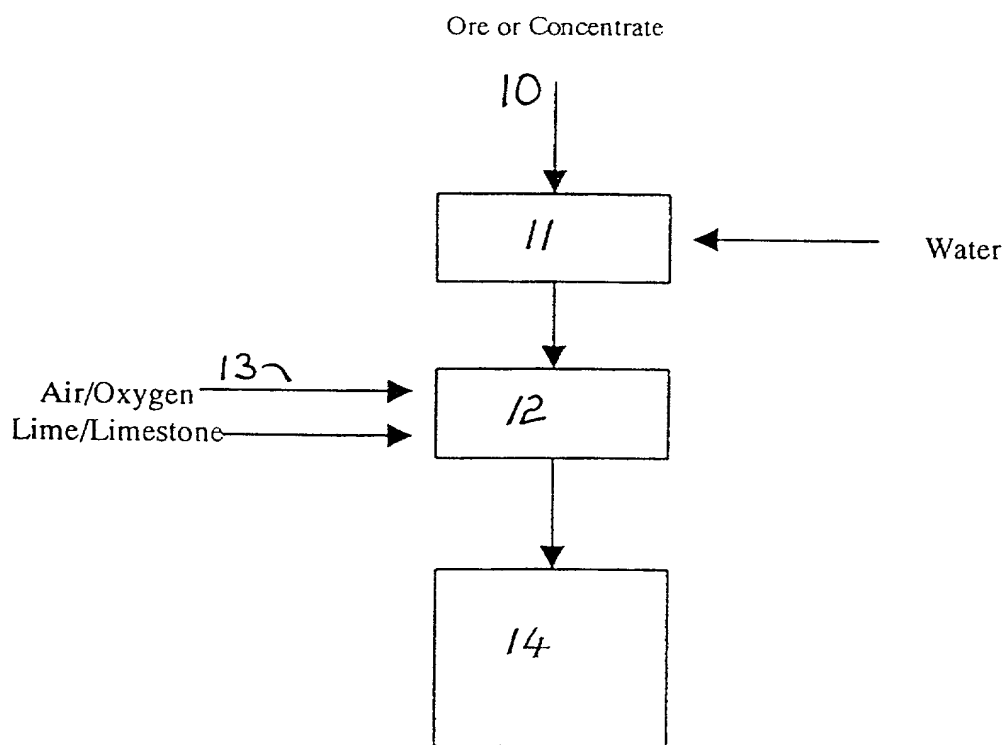


Fig. 1

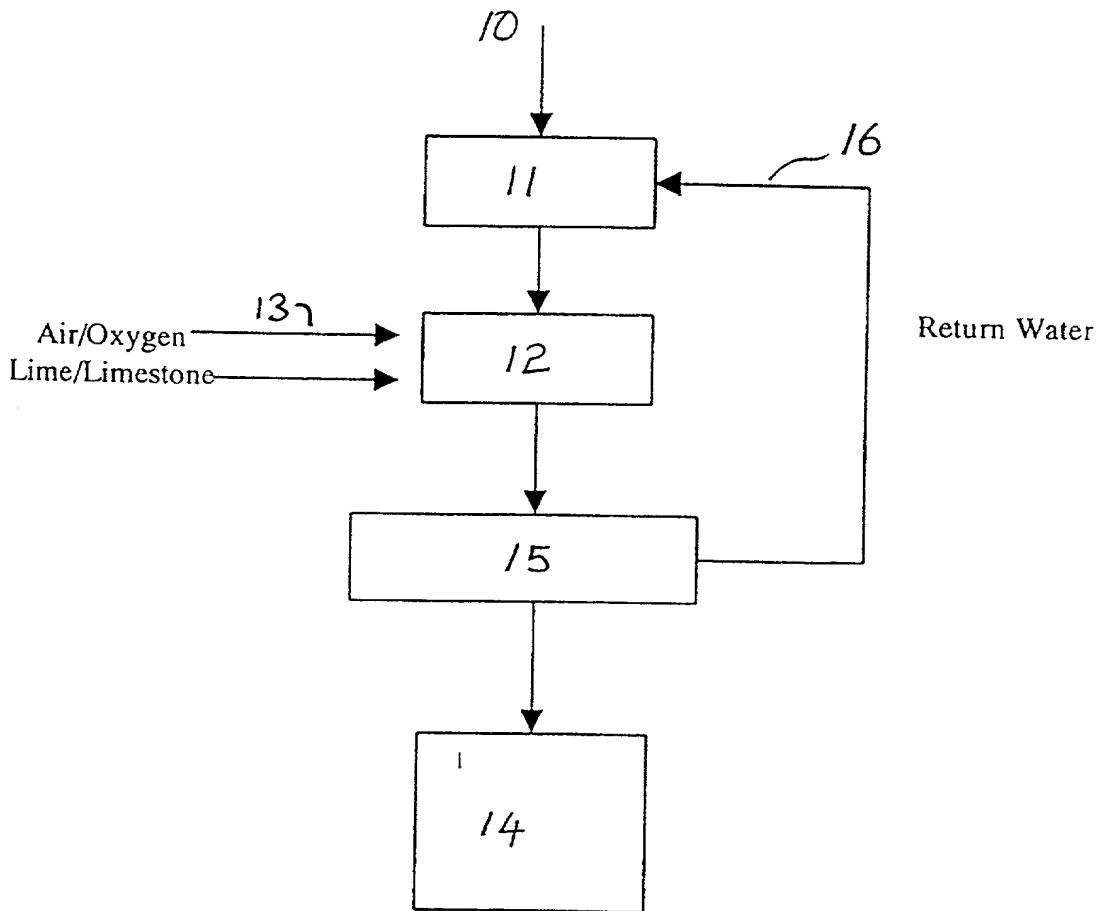


Fig. 2



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US

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As a below named inventor, I hereby declare that:

My residence, post office address, and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Michael Matthew HURN

the specification of which

(Title of the invention)

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Application Number PCT/AU99/00798 was submitted on (MM/DD/YYYY) (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendments specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 356(b) of any foreign application(s) for patent or inventor's certificate, or any PCT International application which designated at least one country other than the United States of America, listed below and have also identified below, by checking this box, any foreign application for patent or inventor's certificate, or of any PCT International application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached?
PP6025	AU	09/21/1998	<input type="checkbox"/>	<input checked="" type="checkbox"/>
PP6313	AU	10/06/1998	<input type="checkbox"/>	<input type="checkbox"/>

☐ Additional foreign application numbers are listed on a supplemental priority data sheet PTO/BB025 attached hereto:

I hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below.

Application Number(s)	Filing Date (MM/DD/YYYY)

☐ Additional provisional application numbers are listed on a supplemental priority data sheet PTO/BB025 attached hereto.

(Page 1 of 2)

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I hereby claim the benefit under 35 U.S.C. 120 of any United States application(s), or 365(c) of any PCT international application designating the United States of America, filed before and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT international application in the manner provided by the first paragraph of 35 U.S.C. 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application.

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Name of Solo or First Inventor: ☐ A petition has been filed for this unsigned inventor

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Residence: City	State
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☐ Additional inventors are being named on the supplemental Additional Inventor(s) sheet(s) PTO/BB/02A attached hereto

(Page 2 of 2)

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## DECLARATION

## ADDITIONAL INVENTOR(S)

Supplemental Sheet

Page 1 of 1

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(July 1998)